PRODUCT INFORMATION



Dexchlorpheniramine (maleate)

Item No. 24027

CAS Registry No.: 2438-32-6

Formal Name: yS-(4-chlorophenyl)-N,N-dimethyl-2-

pyridinepropanamine, 2Z-butenedioate

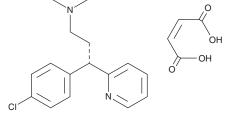
Synonyms: d-Chlorpheniramine (maleate),

S-(+)-Chlorpheniramine (maleate), Fortamine

MF: C₁₆H₁₉CIN₂ • C₄H₄O₄

FW: 390.9 **Purity:** ≥98% Supplied as: A solid Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Laboratory Procedures

Dexchlorpheniramine (maleate) is supplied as a solid. A stock solution may be made by dissolving the dexchlorpheniramine (maleate) in the solvent of choice, which should be purged with an inert gas. Dexchlorpheniramine (maleate) is slightly soluble in chloroform, methanol, DMSO, and dimethyl formamide.

Description

Dexchlorpheniramine is a histamine H₁ receptor antagonist with a pA₂ value of 9.36 in guinea pig ileal tissue in vitro.1 It inhibits the proliferation of previously sensitized, allergen-challenged peripheral blood mononuclear cells by 92% at a concentration of 4.8 μM.² Dexchlorpheniramine reduces noradrenaline uptake in the rat vas deferens ex vivo in response to tyramine stimulation when used at a concentration of 10 μM.3 In vivo, dexchlorpheniramine increases the pain threshold of mice exposed to thermal and chemical stimulation tests when administered intraperitoneally at a dose of 30 mg/kg.⁴ Formulations containing dexchlorpheniramine have been used for the treatment of allergic reactions.

References

- 1. Shamsa, F., Ahmadiani, A., and Khosrokhavar, R. Antihistaminic and anticholinergic activity of barberry fruit (Berberis vulgaris) in the guinea-pig ileum. J. Ethnopharmacol. 64(2), 161-166 (1999).
- Holen, E., Elsayed, S., and Nyfors, A. The effect of H₁ receptor antagonists on peripheral blood mononuclear cells, adenoid cells and primary cell lines. APMIS 103(2), 98-106 (1995).
- Barnett, A., Symchowicz, S., and Taber, R.I. The effects of drugs inhibiting catecholamine uptake on tyramine and noradrenaline-induced contractions of the isolated rat vas deferens. Br. J. Pharmacol. 34(3), 484-492 (1968).
- Farzin, D., Asghari, L., and Nowrouzi, M. Rodent antinociception following acute treatment with different histamine receptor agonists and antagonists. Pharmacol. Biochem. Behav. 72(3), 751-760 (2002).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY

subject to Cayman's Terms and Conditions. Complete Terms and Conditions including Warranty and Limitation of Liability information can be found on our website

Copyright Cayman Chemical Company, 10/03/2022

CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD ANN ARBOR, MI 48108 · USA PHONE: [800] 364-9897

[734] 971-3335

FAX: [734] 971-3640 CUSTSERV@CAYMANCHEM.COM WWW.**CAYMANCHEM**.COM